

**K113165 MINI-MEDIC**Feb 2, 2012  
99 days to decisionK113165 · Product code: **MWI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k113165/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	Oct 26, 2011
Decision date	Feb 2, 2012
Days to decision	99 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Athena Gtx</b>
Location	Des Moines, IA, US
Contact	SEAN MAHONEY
510(k) history	7 submissions · 7 cleared · 2010-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k113165/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 20, 2026